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DOCKET NO.: 48458-0009-00-US 408049 [04-053] Application No.: 10/532.464

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This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

1. (Original) A transgenic organism that produces vanillin when provided with

caffeic acid or an esterified derivative thereof, the organism comprising expressible

transgenes encoding:

a) a 3-O-methyltransferase that catalyzes methylation of caffeic acid to form ferulic

acid; and

b) a chain-shortening enzyme that non-oxidatively converts ferulic acid to vanillin.

2. (Original) The organism of claim 1, which contains an endogenous esterase that

hydrolyzes esters of caffeic acid.

3. (Original) The organism of claim 1, which further comprises an expressible

transgene encoding an esterase that hydrolyzes esters of caffeic acid.

4. (Original) The organism of claim 1, which is a procaryote.

5. (Original) The organism of claim 4, which is Escherichia coli or Pseudomonas

spp.

6. (Original) The organism of claim 1, which is a eucaryote.

7. (Original) The organism of claim 6, which is Pichia pastoris or Saccharomyces

cerevisiae.

8. (Original) The organism of claim 1, wherein the 3-O-methyltransferase is from a

plant source.

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9. (Original) The organism of claim 8. wherein the plant source is selected from the group consisting of Catharanthus roseus, Clarkia breweri, Coffea canephora, Eucalyptus gunnii. Festuca arundinacea, Hordeum vulgare, Lolium perenne, Medicago sativa, Nicotiana tabacum, Ocimum basilicum, Populus tremuloides, Prunus amygdalus, Saccharum officinarum, Sorghum bicolor, Thalictrum tuberosum, Triticum aestivum, Vanilla planifolia and Zea mays.

- 10. (Original) The organism of claim 9, wherein the 3-O-methyltransferase is from Vanilla planifolia.
- (Original) The organism of claim 1, wherein the chain-shortening enzyme is from a plant source.
- (Original) The organism of claim 11, comprising a 4-hydroxybenzaldehyde synthase from Vanilla planifolia.
- 13. (Original) The organism of claim 1. wherein the chain shortening enzyme is from a hacterial source.
 - 14. (Original) The organism of claim 13, comprising enoyl-SCoA hydratase/lyase.
 - 15. (Original) A method for producing vanillin, which comprises:
- a) providing a transgenic organism that produces vanillin when provided with caffeic acid or an esterified derivative thereof, the organism comprising expressible transgenes encoding:
- i) a 3-O-methyltransferase that catalyzes methylation of caffeic acid to form ferulic acid: and
 - ii) a chain-shortening enzyme that non-oxidatively converts ferulic acid to vanillin;
- b) culturing the transgenic organism in the presence of the caffeic acid or esterified derivative thereof, under conditions whereby the transgenic organism produces vanillin; and PHIPM 687156.1

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c) recovering the vanillin from the culture.

16. (Original) The method of claim 15, wherein the organism contains an

endogenous esterase that hydrolyzes esters of caffeic acid.

17. (Original) The method of claim 15, wherein the organism comprises an

expressible transgene encoding an esterase that hydrolyzes esters of caffeic acid.

18. (Original) The method of claim 15, wherein the organism is a procaryote.

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19. (Original) The method of claim 18, wherein the organism is Escherichia coli or

Pseudomonas spp.

20. (Original) The method of claim 15, wherein the organism is a eucaryote.

21. (Original) The method of claim 20, wherein the organism is Pichia pastoris or

Saccharonnyces cerevisiae.

22. (Original) The method of claim 15, wherein the organism comprises a 3-O-

methyltransferase from a plant source.

23. (Original) The method of claim 22, wherein the plant source is selected from the

group consisting of Catharanthus roseus, Clarkia breweri, Coffea canephora, Eucalyptus gunnii. Festuca arundinacea, Hordeum vulgare, Lolium perenne, Medicago sativa, Nicotiana

tabacum, Ocimum basilicum, Populus tremuloides, Prunus amygdalus, Saccharum officinarum, Sorghum bicolor, Thalictrum tuberosum, Triticum aestivum, Vanilla planifolia

and Zea mays.

24. (Original) The method of claim 9, wherein the organism comprises 3-O-

methyltransferase from Vanilla planifolia.

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25. (Original) The method of claim 15, wherein the organism comprises a chain-

26. (Original) The method of claim 25, wherein the organism comprises a 4-

hydroxybenzaldehyde synthase from Vanilla planifolia.

28. (Original) The method of claim 15, wherein the organism comprises a chain

shortening enzyme from a bacterial source.

shortening enzyme from a plant source.

29. (Original) The method of claim 28, wherein the organism comprises enoyl-SCoA

hydratase/lyase.

30. (Original) The method of claim 15, comprising providing the organism with

caffeic acid.

32. (Original) The method of claim 15, comprising providing the organism with a

caffeic acid ester.

33. (Original) The method of claim 33, wherein the caffeic acid ester is one or more

of cichoric acid, rosmarinic acid, chlorogenic acid, 1-caffeolylquinic acid or 1,5-

dicaffeolylquinic acid.

34. (Original) An O-methyltransfcrase from Vanilla planifolia that catalyzes

methylation of substrates selected from the group consisting of 5-OH-ferulic acid ethyl ester,

caffeic acid ethyl ester, caffeoyl aldehyde, 5-OH-coniferaldehyde, 5-OH- ferulic acid. 3,4-

dihydroxybenzaldehyde and caffeic acid.

35. (Original) The O-methyltransferase of claim 34. having an amino acid sequence 5

at least 90% identical to SEQ ID NO:2.

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- 36. (Original) The O-methyltransferase of claim 35, comprising amino acid SEQ ID NO:2.
- 37. (Original) An isolated nucleic acid molecule that encodes the Omethyltransferase of claim 34.
- 38. (Original) The isolated nucleic acid molecule of claim 37, which encodes a polypeptide having an amino acid sequence at least 90% identical to SEQ ID NO:2.
- 39. (Original) The isolated nucleic acid molecule of claim 38, which encodes a polypeptide having SEO ID NO:2.
- (Original) The isolated nucleic acid molecule of claim 39, having a sequence of SEQ ID NO:1.

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